Serial No.: 10/580,202

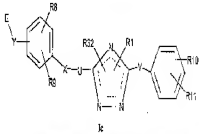
Amendments to the Claims

What is claimed is:

1. (Canceled)

2. (Canceled)

3. (Currently Amended) A compound wherein the compound is of the Formula Ic:



Ic

and stereoisomers or pharmaceutically acceptable salts thereof, and wherein, thereof, wherein:

(a) R1 is hydrogen;

(b) R26, R27, R28 and R29 are each independently selected from the group consisting of hydrogen, hydroxy, oxo, nitro, halo, -SH, $C_1$-$C_4$ alkyl, $C_1$-$C_4$ alkoxy, $CO_2H$, $C_1$-$C_4$ alkoxy, $C_1$-$C_4$ haloalkyl, $C_1$-$C_4$ haloalkyloxy, $C_1$-$C_4$ aminoalkyl, amino, aryl-$C_1$-$C_4$ alkyl, $OC(O)R_{13}$, $CONR_{14}R_{15}$, $OC(O)NR_{16}R_{17}$, $OR_{18}$, $S(O)_2R_{19}$, $NR_{20}R_{21}$, $NR_{22}C(O)R_{23}$, $NR_{24}S(O)_2R_{25}$, $SR_{26}$, $S(O)R_{27}$, $S(O)_2R_{28}$, and $S(O)_2NR_{29}R_{30}$; $R_{13}$, $R_{14}$, $R_{15}$, $R_{16}$, $R_{17}$, $R_{18}$, $R_{19}$, $R_{20}$, $R_{21}$, $R_{22}$, $R_{23}$, $R_{24}$ and $R_{25}$ are each independently selected from the group consisting of hydrogen, $C_1$-$C_6$ alkyl and aryl;

(c) Y is selected from the group consisting of $C_2$-$C_3$ alkyl;

(d) X is selected from the group consisting of a single bond;

(e) U is a divalent linker wherein one carbon atom of the alkylene U chain is substituted with from one to two substituents each independently selected from R30;

2

FILE COVERS 1907 - 30 Oct 2008 VOL 149 ISS 18
FILE LAST UPDATED: 29 Oct 2008 (20081029/ED)

Caplus now includes complete International Patent Classification (IPC) reclassification data for the second quarter of 2008.

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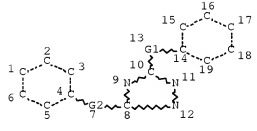

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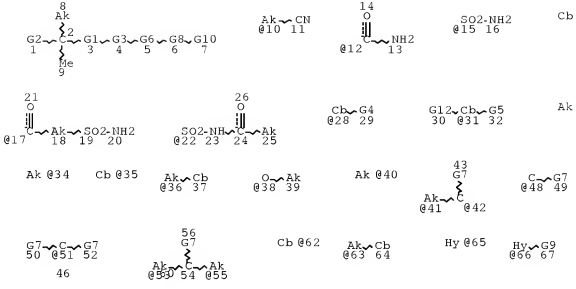

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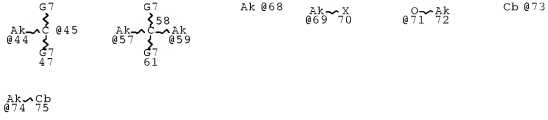

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    ECLA reclassifications to mid August and US national classification
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http://www.stn-international.com/archive/presentations/DWPIAnaVist2_0608.pdf

>>> HELP for European Patent Classifications see HELP ECLA, HELP ICO <<<


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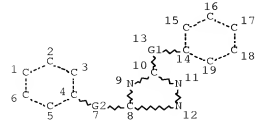

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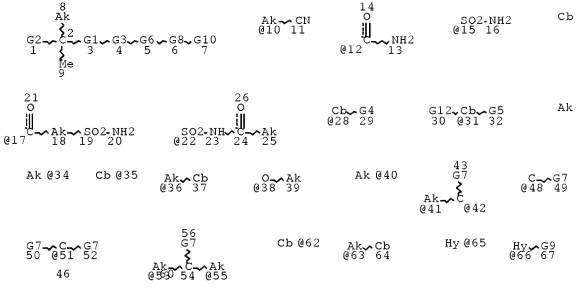


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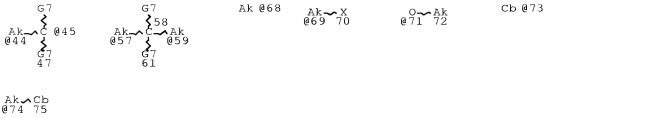


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DOCUMENT NUMBER:          143:153383
TITLE:                    Preparation of triazole, oxadiazole and thiadiazole
                          derivatives as PPAR modulators for the treatment of
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diabetes

INVENTOR(S): Mantlo, Nathan Bryan; Navarro, Antonio; Saeed, Ashraf; Gernert, Douglas Linn; Ma, Tianwei; Pfeifer, Lance Allen

PATENT ASSIGNEE(S): Eli Lilly and Company, USA

SOURCE: PCT Int. Appl., 175 pp.
CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
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| WO 2005065683 | A1 | 20050721 | WO 2004-US39775 | 20041221 |
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OTHER SOURCE(S): CASREACT 143:153383; MARPAT 143:153383

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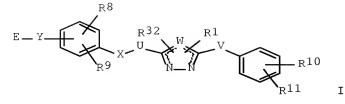

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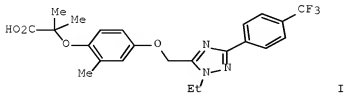

II

AB The title compds. I [X = a single bond, O, S, SO2 and N; U = an aliphatic linker; Y = O, C, S, NH and a single bond; W = N, O or S; E = CR3R4A or A (wherein A = carboxy, tetrazole, alkylnitrile, carboxamide, sulfonamide and acylsufonamide; R3 = H, alkyl, alkoxy; R4 = H, alkyl, alkoxy, etc.; or R3 and R4 are optionally combined to form cycloalkyl); V = (hetero)alkyl, a bond; R1 = H, alkyl, heteroaryl, etc.; R8 = H, alkyl, alkenyl, halo; R9 = H, alkyl, halo, etc.; R10, R11 = H, OH, CN, etc.; R32 = a bond, H, halo, alkyl, etc.] which are modulators of peroxisome proliferator activated receptors (PPARs) and are useful for the treatment of diabetes and other metabolic disorders, were prepared and formulated. E.g., a multi-step synthesis of II, starting from Me glycolate and benzyl bromide, was given. The binding and cotransfection efficacy values for compds. I which are especially useful for modulating a PPAR receptor, are ≤ 100 nM and ≥ 50%, resp.

IT 860260-94-2P 860261-00-3P 860261-28-5P
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RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of triazole, oxadiazole and thiadiazole derivs. as PPAR modulators for the treatment of diabetes)

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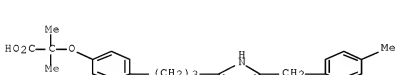

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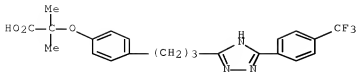

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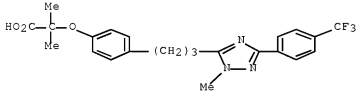

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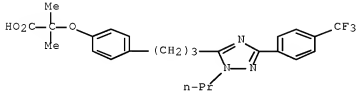

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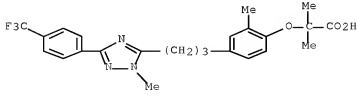

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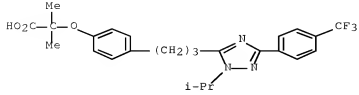

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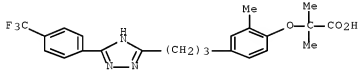
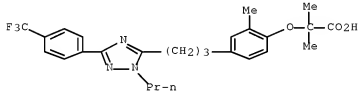
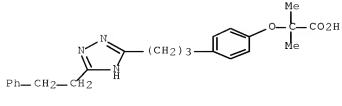


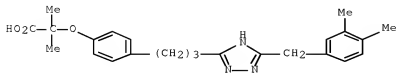
RN 860261-42-3 CAPLUS
CN Propanoic acid, 2-methyl-2-[2-methyl-4-[3-[1-propyl-3-[4-(trifluoromethyl)phenyl]-1H-1,2,4-triazol-5-yl]propyl]phenoxy]- (CA INDEX NAME)



RN 860261-81-0 CAPLUS
CN Propanoic acid, 2-methyl-2-[4-[3-[5-(2-phenylethyl)-1H-1,2,4-triazol-3-yl]propyl]phenoxy]- (CA INDEX NAME)



RN 860262-05-1 CAPLUS
CN Propanoic acid, 2-[4-[3-[5-[(3,4-dimethylphenyl)methyl]-1H-1,2,4-triazol-3-yl]propyl]phenoxy]-2-methyl- (CA INDEX NAME)



REFERENCE COUNT: 10 THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> d his nofil

(FILE 'HOME' ENTERED AT 10:30:07 ON 30 OCT 2008)

FILE 'REGISTRY' ENTERED AT 10:30:16 ON 30 OCT 2008
L1 1 SEA ABB=ON PLU=ON ?METHYLNITRILE?/CNS
D SCA
D
L2 0 SEA ABB=ON PLU=ON ETHYLNITRILE/CNS
ACT YOUNG202/A
---
L3 STR
L4 94763 SEA SSS FUL L3
---
L5 STR
L6 1 SEA SUB=L4 SSS SAM L5
L7 STR L5
L8 1 SEA SUB=L4 SSS SAM L7
L9 10 SEA SUB=L4 SSS FUL L7
L10 0 SEA ABB=ON PLU=ON US200!-580202/APPS

FILE 'CAPLUS' ENTERED AT 11:18:02 ON 30 OCT 2008

FILE 'CAPLUS' ENTERED AT 11:18:06 ON 30 OCT 2008
L11 4 SEA ABB=ON PLU=ON US200!-580202/APPS
L12 1 SEA ABB=ON PLU=ON L11 AND L9
L13 1 SEA ABB=ON PLU=ON L9

FILE 'WPIX' ENTERED AT 11:18:27 ON 30 OCT 2008
L14 0 SEA SSS SAM L7 AND L3
L15 8 SEA SSS FUL L7 AND L3
L16 1 SEA ABB=ON PLU=ON L15/DCR

FILE 'BEILSTEIN' ENTERED AT 11:19:05 ON 30 OCT 2008
L17 0 SEA SSS FUL L7 AND L3

FILE 'MARPAT' ENTERED AT 11:27:15 ON 30 OCT 2008
L18 42 SEA SSS SAM L7
L*** DEL STR
L19 QUE ABB=ON PLU=ON L7

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     FILE 'CAPLUS' ENTERED AT 12:13:20 ON 30 OCT 2008
                D QUE L13

     FILE 'WPIX' ENTERED AT 12:13:29 ON 30 OCT 2008
                D QUE L16

     FILE 'CAPLUS, WPIX' ENTERED AT 12:13:35 ON 30 OCT 2008
L20              1 DUP REM L13 L16 (1 DUPLICATE REMOVED)
                       ANSWER '1' FROM FILE CAPLUS
                D L20 IBIB ABS HITSTR
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